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**From:** "Suplee, Mike"  
**Sent:** Tue 2/12/2008 4:20:02 PM  
**Subject:** Updated agenda for today's meeting 2/12/2008  
Agenda2\_Feb12\_ntrnts.doc

Hi Everyone;

Attached is an updated agenda that is more concise and orders the topic materials better for our discussion today.

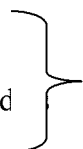
Thanks, Mike

## Updated Agenda for Meeting,, DEQ & EPA Region VIII — Numeric Nutrient Standards

February 12, 2008, 2:00 pm

### Planned Attendance:

Michael Suplee  
Bob Bukantis (may arrive 2:30)  
Tonya Fish  
Ron Steg

1. Update on work & timeframes for numeric nutrient criteria mag, dur, freq (Suplee):
  - a. *Magnitude*-what is driving the numbers I've generated 
    - i. Harm-to-use studies/evaluations
    - ii. Ecoregion-specific stressor-response stud
    - iii. Regional dose-response studies

Reference calibration
  - b. *Duration*-A time period over which exposure is to be averaged
    - i. Seasonal application of nutrient standards
    - ii. Compliance averaging periods
      1. 303(d) listing purposes
      2. Permit compliance
  - c. *Frequency*-How I am determining the proportion of the dataset that can exceed the criteria and still be considered meeting standards
    - i. Consider Figure 1 and our current sampling frames (303d listing)
    - ii. CALM/other states approaches, Clark Fork analysis
2. Timeframe for White Paper (technical basis of criteria) (Suplee)
  - a. Target date for completion
  - b. List of potential academic reviewers
  - c. Assembling/contacting reviewers (Tina)
3. Variance Issue-Can the criteria implementation process be crafted to minimize the need for variances (All)?
  - a. Box 4 of flowchart
  - b. Box 8 & 14 of flowchart
  - c. Box 13 of flowchart
4. Overall timeline to rulemaking (i.e., projecting over next 24 months) (All)
5. Large river nutrient criteria-updates on Yellowstone River project (Suplee)
6. Set next meeting date/time, and draft agenda (All)

Figure 1. Four different stream reaches, each representing 1 sampling frame, showing the variety of sampling sites (hollow dots) that can typically be found along each. Across-time sampling at each dot is possible. We want to consider the selection of statistical tools and their assumptions relative to this sampling reality.

